Toxicological Profiles. This notice solicits public nominations of substances for ATSDR to evaluate for Toxicological Profile development.

DATES: All nominations, whether for substances on the Substance Priority List or for other substances, must be received by August 8, 2022.

ADDRESSES: You may submit nominations, identified by Docket No. ATSDR–2022–0005, by either of the following methods:

- Federal eRulemaking portal at *www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Office of Innovation and Analytics, Agency for Toxic Substances and Disease Registry, 4770 Buford Highway, Mail Stop S102–1, Atlanta, GA 30341–3717. Attn: Docket No. ATSDR–2022–0005.

Instructions: All submissions must include the agency name and docket number for this notice. All relevant comments will be posted without change to *http://www.regulations.gov,* including any personal information provided. Do not submit comments by email. ATSDR does not accept comments by email. This means that no confidential business information or other confidential information should be submitted in response to this notice. Refer to the Submission of Nominations section (below) for the specific information required to be included in a nomination.

FOR FURTHER INFORMATION CONTACT: Kambria Haire, Office of Innovation and Analytics, Agency for Toxic Substances and Disease Registry, 1600 Clifton Rd. NE, Mail Stop S102–1, Atlanta, GA 30329–4027; Email: *ATSDRToxProfileFRNs@cdc.gov;* Phone: 1–800–232–4636.

SUPPLEMENTARY INFORMATION: The Superfund Amendments and Reauthorization Act of 1986 (SARA) [42 U.S.C. 9601 *et seq.*] amended the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA or Superfund) [42 U.S.C. 9601 *et seq.*] by establishing certain requirements for ATSDR and the U.S. Environmental Protection Agency (EPA) concerning hazardous substances most commonly found at facilities on the CERCLA National Priorities List (NPL) (*https://www.epa.gov/superfund/superfund-national-priorities-list-npl*). Among these statutory requirements is a mandate for the Administrator of ATSDR to prepare Toxicological Profiles for each substance included on the Priority List of Hazardous Substances, also known as the Substance Priority list (SPL). This list identifies 275 hazardous substances found at NPL sites that ATSDR and EPA have determined currently pose the most significant potential threat to human health.

Substances To Be Evaluated for Toxicological Profile Development

Each year, ATSDR develops a list of substances to be considered for Toxicological Profile development. The nomination process includes consideration of all substances on ATSDR's SPL, as well as other substances nominated by the public. For more information on ATSDR's SPL, visit *https://www.atsdr.cdc.gov/SPL/.*

Submission of Nominations for Toxicological Profile Development

Today's notice invites voluntary public nominations of substances for toxicological profile development. If nominating a substance that is not on the SPL, please include the rationale for the nomination and any supporting data. ATSDR will evaluate data and information associated with nominated substances and will determine the final list of substances to be chosen for Toxicological Profile development. Substances will be chosen according to ATSDR's specific guidelines for selection. These guidelines can be found in the *Selection Criteria,* which may be accessed at *https://www.atsdr.cdc.gov/toxprofiles/guidance/ATSDR_TP_Selection%20Criteria.pdf.*

Pamela I. Protzel Berman,

Associate Director, Office of Policy, Planning and Partnerships, Agency for Toxic Substances and Disease Registry.

[FR Doc. 2022–14590 Filed 7–7–22; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2022–0014]

Draft Supplemental Environmental Impact Statement; Notice of Public Meeting and Comment Period

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Disease Control and Prevention (CDC) within the Department of Health and Human Services (HHS) announces the opening of a docket and public meeting to obtain comments on the Draft Supplemental Environmental Impact Statement (SEIS) for CDC's Roybal Campus in Atlanta, Georgia. The Draft SEIS was prepared to address changes proposed since completing the 2014 Final Environmental Impact Statement (EIS) for the CDC Roybal Campus 2025 Master Plan (2014 Final EIS) and issuing the Record of Decision dated November 7, 2014. The 2014 Final EIS analyzed the potential impacts associated with implementing a new long-range Master Plan to guide the future physical development of the Roybal Campus for the planning horizon of 2015 to 2025.

DATES: Written comments must be received on or before August 22, 2022.

A virtual public meeting will be held on July 27, 2022, from 6:00 p.m. EST to 8:00 p.m. EST. This meeting will occur via the Zoom platform.

Please register at *https://us06web.zoom.us/meeting/register/tZ0vduiqrT8oEtfyyzvqDUn_oUl5nS-LvfUE.*

Registration is required prior to the meeting. Once registered, you will receive an email with the meeting link and call-in number. The meeting will be recorded using the Zoom platform and a stenographer will transcribe the public meeting. The transcript will be posted on the Docket and included in the Final SEIS.

ADDRESSES: You may submit comments, identified by Docket Number CDC–2022–0014, by either of the following methods:

- *Federal eRulemaking Portal: https://www.regulations.gov/.* Follow the instructions for submitting comments.
- *U.S. Mail:* Thayra Riley, NEPA Coordinator, Office of Safety, Security, and Asset Management, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H20–4, Atlanta, Georgia 30329.

Instructions: All submissions received must include the Agency name and Docket Number (CDC–2022–0014). CDC will post, without change, all relevant comments to *https://www.regulations.gov,* including any personal information provided. *Do not submit comments by email. CDC does not accept comments by email.* For access to the docket to read background documents or comments received, go to *https://www.regulations.gov.* Oral comments on the Draft SEIS will also be accepted during the virtual public meeting scheduled for July 27, 2022.

FOR FURTHER INFORMATION CONTACT: Thayra Riley, NEPA Coordinator, Office of Safety, Security, and Asset Management, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H20–4, Atlanta, Georgia 30329. Email: *cdc-roybalga-*

seis@cdc.gov. Telephone: 770–488–8170.

SUPPLEMENTARY INFORMATION: On January 28, 2022, CDC published a Notice of Intent to prepare an SEIS in the **Federal Register** (87 FR 4603). CDC has prepared a Draft SEIS to analyze the potential impacts of additional proposed components that were not analyzed in the 2014 Final EIS. The proposed components include the addition of a Hospital, Medical, and Infectious Waste Incinerator (HMIWI) in a new laboratory and two emergency standby power diesel generators. The construction of a new laboratory was included in the 2014 Final EIS and will not be re-evaluated in the SEIS.

In accordance with the National Environmental Policy Act (NEPA) as implemented by Council on Environmental Quality (CEQ) regulations (*Code of Federal Regulations* Title 40, Section 1507.3) and HHS environmental procedures, CDC prepared a Draft SEIS to analyze the effects of additional proposed components that were not analyzed in the 2014 Final EIS. The potential impacts of construction and operation of these components on the natural and built environment are being evaluated.

Under NEPA, federal agencies are required to evaluate the environmental effects of their proposed actions and a range of feasible alternatives to the proposed actions prior to making a final decision about what actions to take. The Draft SEIS incorporates the 2014 Final EIS by reference and builds upon that document to focus on specific resource areas that would have potential effects that will differ from those analyzed in the 2014 Final EIS.

Alternatives Considered

CDC analyzed two alternatives in the Draft SEIS: The Proposed Action (Alternative 1) and the No Action Alternative. Alternative 1 consists of the construction and operation of a HMIWI in a new laboratory building and the operation of two emergency standby power diesel generators. The No Action Alternative consists of the construction of the new laboratory without the HMIWI and two emergency standby power generators.

The Draft SEIS evaluates the environmental impacts that may result from Alternative 1 and the No Action Alternative on the following resource categories: air quality, climate change and sustainability, environmental justice, and hazardous/medical/infectious waste. The Draft SEIS identifies measures to mitigate potential adverse impacts.

Availability of the Draft SEIS: Notice of the Availability of the Draft SEIS has been provided to Federal, State, and local agencies and organizations via hard copy letter or electronic mail to the interested parties list. The public is being notified of the availability of the Draft SEIS through this **Federal Register** publication and a notice published in *The Atlanta Journal—Constitution.* The Draft SEIS is available online on the Federal eRulemaking Portal identified by Docket No. CDC–2022–0014. Hard copies of the Draft SEIS are available at the following six locations: Decatur Library, 215 Sycamore Street, Decatur, GA 30030; Toco Hill-Avis G. Williams Library, 1282 McConnell Drive, Decatur, GA 30030; Atlanta-Fulton Public Library, Ponce de Leon Branch, 980 Ponce de Leon Ave. NE, Atlanta, GA 30306; Atlanta-Fulton Public Library, Central Library, One Margaret Mitchell Square, Atlanta, GA 30303; Atlanta-Fulton Public Library, Kirkwood Branch, 11 Kirkwood Rd. NE, Atlanta, GA 30317; and Emory University Robert W. Woodruff Library, 540 Asbury Cir., Atlanta, GA 30322.

Public Meeting: A virtual public meeting will be held on July 27, 2022, from 6:00 p.m. EST to 8:00 p.m. EST. This meeting will occur via the Zoom platform. Please register at *https://us06web.zoom.us/meeting/register/tZ0vduiqrT8oEtfyyzvqDUn_oUl5nS-LvfUE.*

Registration is required prior to the meeting. Once registered, you will receive an email with the meeting link and call-in number.

The meeting will start with a formal presentation and will be followed by a period during which the public can comment or ask questions. A stenographer will transcribe the public meeting. A transcript of the meeting will be made available to the public and will be posted to the public docket at *www.regulations.gov,* identified by Docket No. CDC–2012–0014. CDC will provide a response to comments in the Final SEIS.

Dated: July 1, 2022.

Angela K. Oliver,

Executive Secretary, Centers for Disease Control and Prevention.

[FR Doc. 2022–14518 Filed 7–7–22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; National and Tribal Evaluation of the 2nd Generation of the Health Profession Opportunity Grants (OMB #0970–0462)

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Health Profession Opportunity Grants (HPOG) Program provides healthcare occupational training for Temporary Assistance for Needy Families recipients and other individuals with low incomes. The Office of Management and Budget (OMB) has approved various data collection activities for the National and Tribal Evaluation of the 2nd Generation of HPOG (HPOG 2.0 National and Tribal Evaluation) under OMB #0970–0462. The Administration for Children and Families' (ACF) Office of Planning, Research, and Evaluation (OPRE) is now preparing to conduct the HPOG 2.0 Long-Term Follow-Up Study of HPOG 2.0 participants 5½ years after study enrollment, using a long-term survey (LTS) and administrative data. This notice provides a summary for public review and comment of the use and burden associated with the LTS instrument.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function. You can also obtain copies of the proposed collection of information by emailing *OPREinfocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The HPOG 2.0 evaluation of non-tribal programs is assessing the implementation and impacts of HPOG